

## **EXHIBIT A**

**Puglielli, Maryann**

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**From:** Puglielli, Maryann  
**Sent:** Monday, April 16, 2007 12:52 PM  
**To:** Puglielli, Maryann  
**Subject:** FW: suggestion 10/606229

**Attachments:** prooposal for 10606229 containing allowable subject amtter.doc

> -----Original Message-----

> From: Kim, Vickie  
> Sent: Thursday, December 07, 2006 6:29 PM  
> To: 'carol.einaudi@finnegan.com'  
> Subject: suggestion 10/606229  
>  
> << File: prooposal for 10606229 containing allowable subject amtter.doc >>  
>  
> Vickie Kim  
> Rem. 3A89  
> (571)272-0579  
>  
>  
>

A) I found this information from IDS filed 9/22/06

### What Is In Stieprox Liquid?

Stieprox Liquid contains Ciclopirox Olamine 1.5% w/w as the active ingredient. Ciclopirox Olamine is a broad-spectrum antifungal agent that inhibits the growth of the yeast known which is associated with conditions such as dandruff and seborrheic dermatitis.

Stieprox Liquid also contains sodium laureth sulphate 7.0%, cocamidopropyl betaine, sodium phosphate - dibasic dodecahydrate, citric acid monohydrate, cocamidopropyl betaine, fragrance AF17050 and purified water.

B) I thought Loprox and Stieprox shampoo has same formula as shown in following information (<http://www.pharmacychecker.com/Strength.asp?drugId=58545>) from Google.com :

### ALL STRENGTHS FOR: Loprox Shampoo (called Stieprox IN Canada) - Brand Version

Strength	Click Below For Prices
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1.5 %	<a href="#">View Prices</a>
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C) In specification at page 10:

#### Example 2

A preparation according to the invention has the following composition:

##### Shampoo

5 (based on anionic detergent substance with amphoteric surfactant as cosurfactant)

1-Hydroxy-4-methyl-6-cyclohexyl-2(1H)pyridone	1.00%
Sodium lauryl diglycol ether sulfate (27% strength solution)	36.00%
Cocamidopropylbetaine (30% strength solution)	6.00%
Sodium chloride	3.30%
Water	53.70%

D) In specification at page 6:

anionic, cationic, nonionic or amphoteric nature and can also be present as a combination of these substances.

Preferably, however, anionic surfactants are employed on their own or as a 5 mixture with other anionic surfactants as base surfactants - if appropriate with addition of amphoteric surfactants as cosurfactant.

As the sole detergent substances, amphoteric surfactants are virtually 10 insignificant, since their foaming behavior, thickenability and partly also skin and eye mucous membrane tolerability are only moderate. In combination with various anionic surfactants, however, precisely these properties are synergistically improved. This explains the relatively great importance of the amphoteric surfactants for the optimization of anionic shampoo-bases.

E) your remarks: Loprox has unexpected commercial success(see your remarks at page 13)

Application No. 10/606,229  
Attorney Docket No. 03804.1596-01

continued growth in sales has occurred despite the presence in the market of competitive, mostly less expensive, prescription products. *Id.*, Paragraphs 6 and 7.

This 13% increase in market share in an adverse market is an indicator of commercial success and would not have occurred unless physicians and patients judged Loprox Shampoo, even at a premium price, to be a more effective treatment for seborrheic dermatitis than the competitive prescription shampoos already on the market. *Id.*

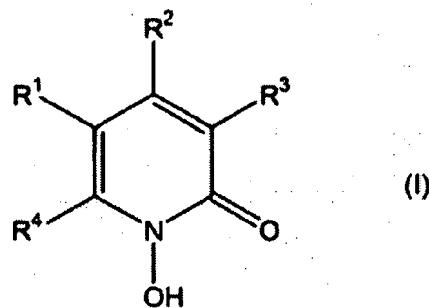
My proposal is based on combination of these factors shown above.

Here's my proposal.

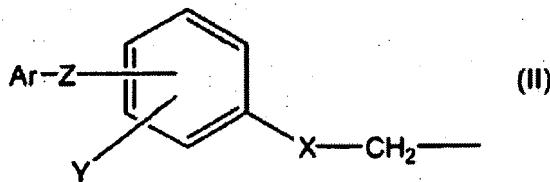
Claim 14 (currently amended)

14. (Currently Amended) A method of treating seborrheic dermatitis comprising administering to a human seborrheic dermatitis patient an amount effective for the treatment of seborrheic dermatitis of a composition comprising:

(A) [[an]] a sole active component consisting of at least one 1-hydroxy-2-pyridone of formula I, wherein the at least one 1-hydroxy-2-pyridone is present in free form or as a pharmaceutically acceptable salt:



where  $R^1$ ,  $R^2$ , and  $R^3$ , which are identical or different, are H or alkyl having 1 to 4 carbon atoms, and  $R^4$  is a saturated hydrocarbon radical having 6 to 9 carbon atoms or a radical of formula II:



where:

X is S or O;

Y is H, or 1 or 2 identical halogen atoms, or a mixture of 2 different halogen atoms;

Z is a single bond, or a linking radical comprising

(1) O, or

(2) S, or

(3)  $-\text{CR}_2-$ , where R is H or  $(\text{C}_1\text{-C}_4)$ -alkyl, or

(4) from 2 to 10 carbon atoms linked in the form of a straight or branched chain, which optionally further comprises one or more of the following:

(i) carbon-carbon double bond, and

(ii) O, S, or a mixture thereof, wherein if 2 or more O or S atoms or a mixture thereof are present, each O or S atom is separated by at least 2 carbon atoms; and,

In any of the foregoing linking radicals, any remaining free valences of the carbon atoms of said linking radical are saturated by H, (C<sub>1</sub>-C<sub>4</sub>)-alkyl, or a mixture thereof;

and

Ar is an aromatic ring system having one or two rings, the aromatic ring system being unsubstituted or substituted by one, two, or three radicals, which are

identical or different, and are chosen from halogen, methoxy, (C<sub>1</sub>-C<sub>4</sub>)-alkyl, trifluoromethyl, and trifluoromethoxy; and

(B) an anionic surfactant as a base surfactant with amphoteric surfactant as cosurfactant; wherein the composition has a pH ranging from about 4.5 to about 6.5; and wherein the composition is in the form of shampoo and the said 1-hydroxy-2-pyridone is 1-hydroxy-4-methyl-6-cyclohexyl-2(1H) pyridine.

## **EXHIBIT B**

## LOPROX® SHAMPOO (ciclopirox) 1%

**Rx Only**  
**FOR TOPICAL USE ONLY**  
**NOT FOR OPHTHALMIC, ORAL OR INTRAVAGINAL USE**  
**KEEP OUT OF REACH OF CHILDREN**

### DESCRIPTION

LOPROX® (ciclopirox) Shampoo 1% contains the synthetic antifungal agent, ciclopirox. Each gram (equivalent to 0.96 mL) of LOPROX Shampoo contains 10 mg ciclopirox in a shampoo base consisting of Purified Water USP, Sodium Laureth Sulfate, Disodium Laureth Sulfosuccinate, Sodium Chloride USP, and Laureth-2. LOPROX Shampoo is a colorless, translucent solution. The chemical name for ciclopirox is 6-cyclohexyl-1-hydroxy-4-methyl-2(1H)-pyridone, with the empirical formula  $C_{12}H_{11}NO_2$  and a molecular weight of 207.27. The CAS Registry Number is [29342-05-0]. The chemical structure is:



### CLINICAL PHARMACOLOGY

#### Mechanism of Action

Ciclopirox is a hydroxypyridone antifungal agent although the relevance of this property for the indication of seborrheic dermatitis is not known. Ciclopirox acts by chelation of polyvalent cations ( $Fe^{3+}$  or  $Al^{3+}$ ), resulting in the inhibition of the metal-dependent enzymes that are responsible for the degradation of peroxides within the fungal cell.

#### Pharmacokinetics and Pharmacodynamics

In a study in patients with seborrheic dermatitis of the scalp, application of 5 mL ciclopirox shampoo 1% twice weekly for 4 weeks, with an exposure time of 3 minutes per application, resulted in detectable serum concentrations of ciclopirox in 6 out of 18 patients. The serum concentrations measured throughout the dosing interval on Days 1 and 29 ranged from 10.3 ng/mL to 13.2 ng/mL. Total urinary excretion of ciclopirox was less than 0.5% of the administered dose.

### CLINICAL STUDIES

In two randomized, double-blind clinical trials, patients 16 years and older with seborrheic dermatitis of the scalp applied LOPROX Shampoo or its vehicle two times per week for 4 weeks. Patients who were immunocompromised, those with psoriasis or atopic dermatitis, women of childbearing potential not using adequate contraception, and pregnant or lactating women were excluded from the clinical studies. An evaluation of the overall status of the seborrheic dermatitis, and the presence and severity of erythema or inflammation, and scaling, was made at week 4, using a scale of 0 = none, 1 = slight, 2 = mild, 3 = moderate, 4 = pronounced, and 5 = severe. Effective treatment was defined as achieving a score of 0 (or a score of 1 if the baseline score was 2-3) simultaneously for status of the seborrheic dermatitis, erythema or inflammation, and scaling at Week 4. Ciclopirox shampoo was shown to be statistically significantly more effective than vehicle in both studies. Efficacy results for the two studies are presented in the following table.

#### Effective Treatment Rates at Week 4 in Studies 1 and 2

	Ciclopirox Shampoo	Vehicle
Study 1	220/380 (58%)	60/192 (31%)
Study 2	65/250 (26%)	32/249 (13%)

Efficacy for black patients was not demonstrated, although only 53 black patients were enrolled in the two pivotal studies.

#### Microbiology

Ciclopirox is fungicidal *in vitro* against *Malassezia furfur* (*Pityrosporum spp.*), *P. ovale*, and *P. orbiculare*. Ciclopirox acts by chelation of polyvalent cations ( $Fe^{3+}$  or  $Al^{3+}$ ), resulting in the inhibition of the metal-dependent enzymes that are responsible for the degradation of peroxides within the fungal cell.

The clinical significance of antifungal activity in the treatment of seborrheic dermatitis is not known.

### INDICATIONS AND USAGE

LOPROX Shampoo is indicated for the topical treatment of seborrheic dermatitis of the scalp in adults.

### CONTRAINDICATIONS

LOPROX Shampoo is contraindicated in individuals who have shown hypersensitivity to any of its components.

### WARNINGS

LOPROX Shampoo is not for ophthalmic, oral, or intravaginal use.

Keep out of reach of children.

### PRECAUTIONS

#### General

If a reaction suggesting sensitivity or irritation should occur with the use of LOPROX Shampoo, treatment should be discontinued and appropriate therapy instituted.

Contact of LOPROX Shampoo with the eyes should be avoided. If contact occurs, rinse thoroughly with water.

Seborrheic dermatitis may appear at puberty, however, no clinical studies have been done in patients younger than 16 years.

There is no relevant clinical experience with patients who have a history of immunosuppression (e.g., extensive, persistent, or unusual distribution of dermatomycoses, recent or recurring herpes zoster, or persistent herpes simplex), who are immunocompromised (e.g., HIV-infected patients and transplant patients), or who have a diabetic neuropathy.

#### **Information for Patients**

The patient should be instructed to:

1. Use LOPROX Shampoo as directed by the physician. Avoid contact with the eyes and mucous membranes. If contact occurs, rinse thoroughly with water. LOPROX Shampoo is for external use on the scalp only. Do not swallow.
2. Use the medication for seborrheic dermatitis for the full treatment time even though symptoms may have improved. Notify the physician if there is no improvement after 4 weeks.
3. Inform the physician if the area of application shows signs of increased irritation (redness, itching, burning, blistering, swelling, or oozing) indicative of possible allergic reaction.
4. Not use the medication for any disorder other than that for which it is prescribed.

#### **Carcinogenesis, Mutagenesis, and Impairment of Fertility:**

Long-term animal studies have not been performed to evaluate the carcinogenic potential of LOPROX Shampoo or ciclopirox. The following *in vitro* genotoxicity tests have been conducted with ciclopirox: evaluation of gene mutation in the Ames *Salmonella* and *E. coli* assays (negative); chromosome aberration assays in V79 Chinese hamster lung fibroblast cells, with and without metabolic activation (positive); chromosome aberration assays in V79 Chinese hamster lung fibroblast cells in the presence of supplemental Fe<sup>2+</sup>, with and without metabolic activation (negative); gene mutation assays in the HGPRT test with V79 Chinese hamster lung fibroblast cells (negative); and a primary DNA damage assay (i.e., unscheduled DNA synthesis assay in A549 human cells) (negative). An *in vitro* cell transformation assay in BALB/c 3T3 cells was negative for cell transformation. In an *in vivo* Chinese hamster bone marrow cytogenetic assay, ciclopirox was negative for chromosome aberrations at a dosage of 5000 mg/kg body weight.

A combined oral fertility and embryofetal developmental study was conducted in rats with ciclopirox olamine. No effect on fertility or reproductive performance was noted at the highest dose tested of 3.85 mg/kg/day ciclopirox (approximately 1.3 times the maximum recommended human dose based on body surface area comparisons).

#### **Pregnancy:**

##### **Teratogenic effects: Pregnancy Category B**

Oral embryofetal developmental studies were conducted in mice, rats, rabbits and monkeys. Ciclopirox or ciclopirox olamine was orally administered during the period of organogenesis. No maternal toxicity, embryotoxicity or teratogenicity were noted at the highest doses of 77, 125, 80 and 38.5 mg/kg/day ciclopirox in mice, rats, rabbits and monkeys, respectively (approximately 13, 42, 54 and 26 times the maximum recommended human dose based on body surface area comparisons, respectively).

Dermal embryofetal developmental studies were conducted in rats and rabbits with ciclopirox olamine dissolved in PEG 400. Ciclopirox olamine was topically administered during the period of organogenesis. No maternal toxicity, embryotoxicity or teratogenicity were noted at the highest doses of 92 mg/kg/day and 77 mg/kg/day ciclopirox in rats and rabbits, respectively (approximately 31 and 54 times the maximum recommended human dose based on body surface area comparisons, respectively).

There are no adequate or well-controlled studies of topically applied ciclopirox in pregnant women. Because animal reproduction studies are not always predictive of human response, LOPROX Shampoo should be used during pregnancy only if clearly needed.

#### **Nursing Mothers:**

It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when LOPROX Shampoo is administered to a nursing woman.

#### **Pediatric Use:**

Seborrheic dermatitis may appear at puberty, however, no clinical studies have been done in patients younger than 16 years.

#### **Geriatric Use:**

In clinical studies, the safety and tolerability of LOPROX Shampoo in the population 65 years and older was comparable to that of younger subjects. Results of the efficacy analysis in those patients 65 years and older showed effectiveness in 25 of 85 (29%) patients treated with LOPROX Shampoo, and in 15 of 61 (25%) patients treated with the vehicle; due to the small sample size, a statistically significant difference was not demonstrated. Other reported clinical experience has not identified differences in responses between the elderly and younger subjects, but greater sensitivity to adverse effects in some older individuals cannot be ruled out.

#### **ADVERSE REACTIONS**

In 626 patients treated with LOPROX Shampoo twice weekly in the two pivotal clinical studies, the most frequent adverse events were increased itching in 1% of patients, and application site reactions, such as burning, erythema, and itching, also in 1% of patients. Other adverse events occurred in individual patients only.

Adverse events that led to early study medication termination in clinical trials occurred in 1.5% (26/1738) of patients treated with LOPROX Shampoo and 2.0% (12/661) of patients treated with shampoo vehicle. The most common adverse events leading to termination of study medication in either group was seborrhea. In the LOPROX Shampoo group, other adverse events included rash, pruritis, headache, ventricular tachycardia, and skin disorder. In the shampoo vehicle group, other adverse events included skin disorder and rash.

#### **DOSAGE AND ADMINISTRATION**

Wet hair and apply approximately 1 teaspoon (5 mL) of LOPROX Shampoo to the scalp. Up to 2 teaspoons (10 mL) may be used for long hair. Lather and leave on hair and scalp for 3 minutes. A timer may be used. Avoid contact with eyes. Rinse off. Treatment should be repeated twice per week for 4 weeks, with a minimum of 3 days between applications.

If a patient with seborrheic dermatitis shows no clinical improvement after 4 weeks of treatment with LOPROX Shampoo, the diagnosis should be reviewed.

#### **HOW SUPPLIED**

LOPROX® (ciclopirox) Shampoo 1% is supplied in 120 mL plastic bottles (NDC 99207-010-10). Discard unused product after initial treatment duration. Store between 15°C and 30°C (59°F and 86°F).

#### **Manufactured for:**

**MEDICIS® Pharmaceutical Corp.**

**Scottsdale, AZ 85258**

**by: Patheon, Inc.**

**Mississauga, Ontario LSN 7K9**

**CANADA**